

Dibenz[a,h]anthracene

Division of Safety National Institutes of Health



WARNING!

THIS COMPOUND IS ABSORBED THROUGH THE SKIN AND RESPIRATORY AND INTESTINAL TRACTS. IT IS CARCINOGENIC AND MAY IRRITATE TISSUES AND INDUCE SENSITIVITY. AVOID FORMATION AND BREATHING OF DUSTS.

LABORATORY OPERATIONS SHOULD BE CONDUCTED IN A FUME HOOD, GLOVE BOX, OR VENTILATED CABINET.

AVOID SKIN CONTACT: IF EXPOSED, WASH WITH SOAP AND WATER. AVOID WASHING WITH SOLVENTS AND EXPOSURE TO UV LIGHT.

FOR EYE EXPOSURE, IRRIGATE IMMEDIATELY WITH LARGE AMOUNTS OF WATER. FOR INGESTION, INDUCE VOMITING. FOR INHALATION, REMOVE VICTIM PROMPTLY TO CLEAN AIR. ADMINISTER RESCUE BREATHING IF NECESSARY. REFER TO PHYSICIAN.

IN CASE OF LABORATORY SPILL, WEAR PROTECTIVE CLOTHING DURING CLEANUP. AVOID SKIN CONTACT OR BREATHING OF DUST. USE ORGANIC SOLVENT (NOT ALCOHOL) TO DISSOLVE COMPOUND. WASH DOWN AREA WITH SOAP AND WATER. CHECK FOR FLUORESCENCE OF RESIDUES WITH UV LIGHT. DISPOSE OF WASTE SOLUTIONS AND MATERIALS BY INCINERATION.

A. Background

Dibenz[a,h]anthracene (DBA) is well established as a carcinogen, intermediate in activity between benz[a]anthracene and 7,12-dimethylbenz[a]anthracene. DBA is detected as an environmental contaminant at levels generally lower than benzo[a]pyrene or benz[a]anthracene. It has no known commercial or industrial use and is employed solely in carcinogenesis research. It is destroyed through photooxidation in the atmosphere and is believed to be degraded slowly by bacteria in the soil.

issued 11/81

Chemical and Physical Data 1. Chemical Abstract No.: 53-70-3

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Vapor pressure:

- 2. Synonyms:
 - DBA 1.2.5.6-Dibenzanthracene
 - DB(a,h)A 1,2:5,6-Dibenzanthracene 1.2.5.6-Dibenzanthracene
 - Molecular formula: structure: C22H14 weight: 278.33
- Density: 1.282 g/cm^3 . 4. Absorption spectroscopy: UV (Friedel and Orchin, 1951; Pe kamper et al., 1967); UV fluorescence (Sawicki et al., 196
- 5. IR (Sadtler, 1961; Pouchert, 1970); NMR (Clin and Lemancea 1970: Haigh et al., 1970: Ozubko et al., 1974).

No data.

- 7. Solubility: Soluble in most organic solvents; slightly so in alcohol; solubility in water, 0.0005 mg/liter at 27°C. Description, appearance: Colorless plates. 8.
- 9. Boiling point: 524°C. Melting point: 267°C.
- 10. Stability: Stable in dark at ambient temperature or below Solutions undergo photooxidation in air and light.
- Chemical reactivity: Not spontaneously reactive, but ente 11.
- into numerous types of reactions with organic reagents.

Does not apply.

12. Flash point: Does not apply.

Flammable limits:

13. Autoignition temperature: No data.

In powdered

- DBA does not require special fire-fighting procedures or equip-1.
- ment. Because of the electrostatic nature of dry DBA, fire
- fighters should wear full-face masks.
- DBA does not present unusual fire and explosion hazards. 2.
- 3. DBA is unstable in presence of light and is more unstable when UV radiation is present.
- 4. Incompatibilities:
- No data.
- 5. DBA is not known to produce hazardous decomposition products.
- DBA is nonvolatile and does not require nonspark equipment. 6.
- When handled in flammable solvents such as benzene, the pre-
- cautions required for such solvents will apply.
- form DBA is electrostatic, and when used in this form, it requires the use of antistatic devices.
- Operational Procedures The NIH Guidelines for the Laboratory Use of Chemical Carcinogens
- describe operational practices to be followed when potentially carcinogenic chemicals are used in NIH laboratories. lines should be consulted to identify the proper use conditions
- required and specific controls to be implemented during normal and
- complex operations or manipulations involving DBA.
 - 1. Chemical inactivation: No validated method reported.
 - 2. Decontamination: Turn off equipment that could be affected by DBA or the materials used for cleanup. If more than 1 g
 - has been spilled or if there is any uncertainty regarding the procedures to be followed for decontamination, call the NIH
 - Fire Department (dial 116) for assistance. Wash surfaces with copious quantities of soap and water. Glassware should be rinsed (in a hood) with an organic solvent (not alcohol), followed by soap and water. Animal cages should be washed with
 - soap and water. 3. No waste streams containing DBA shall be disposed
 - of in sinks or general refuse. Surplus DBA or chemical waste streams contaminated with DBA shall be handled as hazardous chemical waste and disposed of in accordance with the NIH chemical waste disposal system. Nonchemical waste (e.g.,

disinfected by heat using a standard autoclave treatment and packaged for incineration, as above. Burnable waste (e.g., absorbent bench top liners) minimally contaminated with DBA shall be handled as potentially infectious waste and packaged

animal carcasses and bedding) containing DBA shall be handled and packaged for incineration in accordance with the NIH

medical-pathological waste disposal system. Potentially infectious waste (e.g., tissue cultures) containing DBA shall be

for incineration, as above. Absorbent materials (e.g., associated with spill cleanup) grossly contaminated shall be

handled in accordance with the chemical waste disposal system. Radioactive waste containing DBA shall be handled in accordance

4. Storage: Store solid DBA and its solutions in dark-colored. tightly closed containers, preferably under refrigeration.

with the NIH radioactive waste disposal system.

- Monitoring and Measurement Procedures Including Direct Field Measurements and Sampling for Subsequent Laboratory Analysis (Jones and Freudenthal, 1978)
 - 1. Sampling: Two methods are recommended: using an adsorption sampler in which cooled air is passed through Tenax and using
 - high-volume filtration through fiberglass filter traps. Separation and analysis: Several methods are available and
- 2. offer various degrees of sensitivity. For separation, TLC, HPLC, and GC are useful. TLC is the least efficient of these three methods. HPLC and GC are highly efficient. The most
- useful and sensitive method for separation and analysis of DBA is GC-MS. This method allows for accurate identification in the nanogram to picogram level; it is still desirable to confirm the identification by other analytical methods. UV spectroscopy is useful but is limited because of possible

similarity in spectra with a related compound. Fluorescence

cyclic aromatic hydrocarbons it may be assumed that DBA should be readily absorbed through the skin, by intravenous and intra

- spectroscopy gives both excitation and emission spectra and its sensitivity level is in the nanogram range. It is more sensitive than UV by a factor of 10^2 or 10^3 or greater. methods are phosphorescence, NMR, and IR spectroscopy.
- Biological Data (Animal and Human)
 - Absorption: There are no data, but in analogy with other poly 1.

peritoneal injection, and by inhalation.

containing tissues. Metabolism and excretion: Like other polycyclic aromatic hydr 3.

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- carbons, DBA is metabolized by the aryl hydrocarbon hydroxylas system of mammalian liver to a variety of epoxides, diols, and phenols, some of which are responsible for the toxic and carcinogenic action of DBA (Kuroki et al., 1972). There are no
- data on excretion products but they are likely to be conjugate of these metabolites. 4. Toxic effects: There are no data on the acute toxicity of DBA As a class, polycyclic aromatic hydrocarbons are regarded as having low acute toxicity in animals and man (Boyland et al.,

Distribution: No data, but probably distributed to lipid-

1965; Heidelberger, 1975). There is no specific target organ but rather a general toxic (and carcinogenic) effect on epithelial and fibroblastic cells. Carcinogenic effects: DBA is moderately carcinogenic in exper-5. mental animals. Lung and skin carcinomas have been reported in mice after parenteral or topical treatment with DBA, and fibro-

sarcomas have been produced in rats after subcutaneous injec-

6. Mutagenic and teratogenic effects: DBA is mutagenic in the Ames test and in mammalian cell cultures in the presence of a metabolizing system. There is no evidence for teratogenicity.

Emergency Treatment

tion.

- Skin and eye exposure: For skin exposure, remove contaminated 1. clothing and wash skin with soap and water. Skin should not be rinsed with organic solvents or scanned with UV light. For
- eye exposure, irrigate immediately with copious quantities of
- running water for at least 15 minutes.
- Ingestion: Drink plenty of water. Induce vomiting. 2.
- 3. Inhalation: Remove victim promptly to clean air. Administer rescue breathing if necessary. 4. Refer to physician.

References Boyland, E., P. Sims, and C. Huggins. 1965. Induction of adrenal damage and cancer with metabolites of 7,12-dimethylbenz[a]anthracene. Nature 207:816-817.

Raven Press, New York.
 Ruroki, T., E. Huberman, H. Marquardt, J.K. Selkirk, C. Heidelberger P.L. Grover, and P. Sims. 1972. Binding of K-region epoxides and other derivatives of benz[a]anthracene and dibenz[a,h]anthracene to DNA, RNA, and proteins of transformable cells. Chem-Biol Interact 4:389-397.
 Ozubko, R.S., G.W. Buchnan, and I.C. Smith. 1974. Carbon-13 nuclea

Jin, B., and B. Lemanceau. 1970. Nuclear magnetic resonance study of carcinogenic and non-carcinogenic isomers (dibenzacridines and dibenzanthracenes). CR Acad Sci (Paris), Ser D 271:788-

riedel, R.A., and M. Orchin. 1951. Ultraviolet Spectra of Aro-

laigh, C.W., R.B. Mallion, and E.A. Armour. 1970. Proton magnetic resonance of planar condensed benzenoid hydrocarbons. II. A critical evaluation of the McWeeny "ring current" theory.

leidelberger, C. 1975. Chemical carcinogenesis. Annu Rev Biochem

lones, P.W., and R.I. Freudenthal, eds. 1978. Carcinogenesis, Vol.

matic Compounds. John Wiley and Sons, New York.

790.

44:79-121.

Molec Phys 18:751-766.

- Chem-Biol Interact 4:389-397.

 Ozubko, R.S., G.W. Buchnan, and I.C. Smith. 1974. Carbon-13 nuclea magnetic resonance spectra of carcinogenic polynuclear hydrocarbons. I. 3-Methyl-cholanthrene and related benzanthracenes Can J Chem 52:2493-2501.

 Overkamper H.H. I. Sandeman and C.J. Timmores eds. 1967. DMS III
- Can J Chem 52:2493-2501.

 Perkamper, H.H., I. Sandeman, and C.J. Timmores, eds. 1967. DMS UNATION OF Compounds, Vol. III. Weinheim: Verlag Chemic Butterworths, E5/6, London.
- Butterworths, E5/6, London.

 Pouchert, C.E., ed. 1970. Aldrich Library of Infrared Spectra.
 Aldrich Chemical Co., Inc., Milwaukee, WI.
- Sadtler Standard Spectra. 1961. Sadtler Research Laboratories,
 Inc., Philadelphia, PA.
- Sawicki, E., T.R. Hauser, and T.W. Stanley. 1960. Ultraviolet visible and fluorescence spectral analysis of polynuclear hydrocarbons. Int J Air Pollution 2:253-276.